# LORATADINE- loratadine tablet Cabinet Health P.B.C.

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Loratadine

### **Drug Facts**

### Active ingredient (in each tablet)

Loratadine USP 10 mg

#### **Purpose**

**Antihistamine** 

#### Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

• runny nose • itchy, watery eyes • sneezing • itching of the nose or throat

### Warnings

#### Do not use

if you have ever had an allergic reaction to this product or any of its ingredients.

## Ask a doctor before use if you have

liver or kidney disease. Your doctor should determine if you need a different dose.

## When using this product

do not take more than directed. Taking more than directed may cause drowsiness.

## Stop use and ask a doctor if

an allergic reaction to this product occurs. Seek medical help right away.

## If pregnant or breast-feeding,

ask a health professional before use.

## Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

#### **Directions**

adults and children 6 years and over 1 tablet daily; not more than 1 tablet in 24 hours children under 6 years of age ask a doctor consumers with liver or kidney disease ask a doctor

#### Other information

- Tamper-evident: do not use if foil seal under cap, printed with "SEALED for YOUR PROTECTION" is missing, open or broken
- store between 20° to 25°C (68° to 77°F)

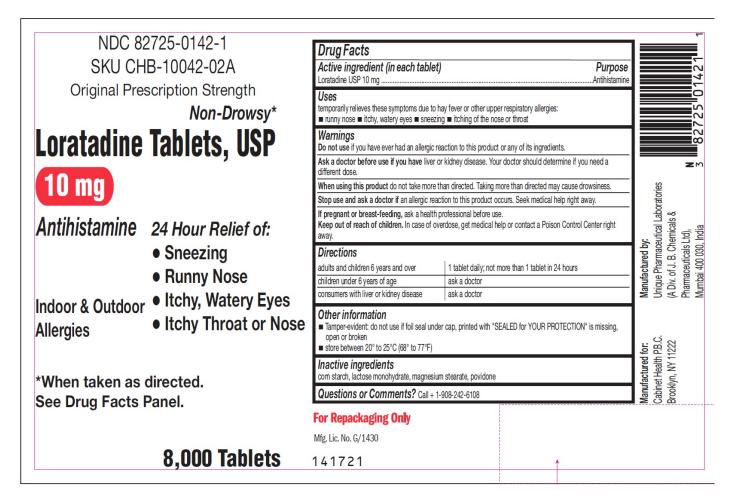
#### Inactive ingredients

corn starch, lactose monohydrate, magnesium stearate, povidone

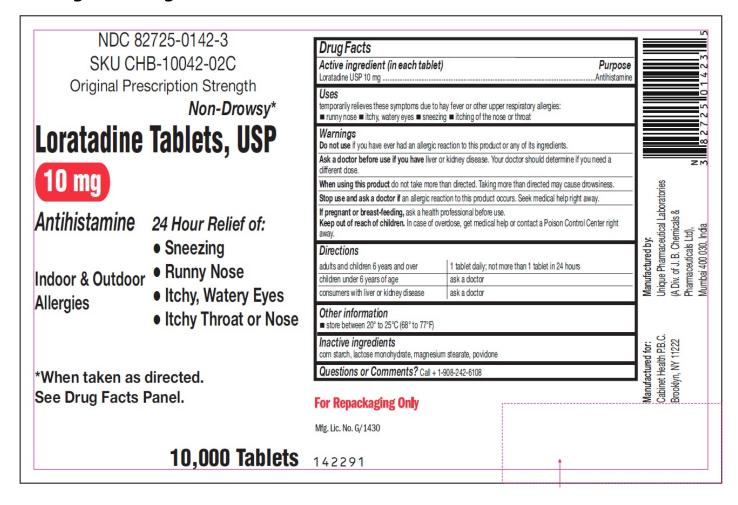
#### Questions or Comments?

Call + 1-908-242-6108

Package Labeling: 82725-0142-1



## Package Labeling: 82725-0142-3



Package Labeling: 82725-0142-4

NDC 82725-0142-4 **Drug Facts** SKU CHB-10042-02D Active ingredient (in each tablet) Purpose Loratadine USP 10 mg .. .Antihistamine Original Prescription Strength Uses temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: Non-Drowsy\* ■ runny nose ■ itchy, watery eyes ■ sneezing ■ itching of the nose or throat **Loratadine Tablets, USP** Do not use if you have ever had an allergic reaction to this product or any of its ingredients. Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a 10 mg When using this product do not take more than directed. Taking more than directed may cause drowsiness. Unique Pharmaceutical Laboratories Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away. If pregnant or breast-feeding, ask a health professional before use Antihistamine 24 Hour Relief of: (A Div. of J. B. Chemicals & Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right Pharmaceuticals Ltd), Mumbai 400 030, India Sneezing Manufactured by: Directions Runny Nose adults and children 6 years and over 1 tablet daily; not more than 1 tablet in 24 hours Indoor & Outdoor • Itchy, Watery Eyes children under 6 years of age ask a doctor consumers with liver or kidney dise **Allergies** Other information Itchy Throat or Nose ■ store between 20° to 25°C (68° to 77°F) Inactive ingredients Manufactured for: Cabinet Health P.B.C. Brooklyn, NY 11222 Questions or Comments? Call + 1-908-242-6108 \*When taken as directed. For Repackaging Only See Drug Facts Panel. Mfg. Lic. No. G/1430

142292

Package Labeling: 82725-0142-2

**96,000 Tablets** 

NDC 82725-0142-2 SKU CHB-10042-02B

Original Prescription Strength

Non-Drowsy\*

# **Loratadine Tablets, USP**



## Antihistamine 24 Hour Relief of:

- Sneezing
- Runny Nose

Indoor & Outdoor **Allergies** 

- Itchy, Watery Eyes
- Itchy Throat or Nose

\*When taken as directed. See Drug Facts Panel.

#### **Drug Facts** Active ingredient (in each tablet) Purpose Loratadine USP 10 mg Antihistamin temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: ■ runny nose ■ itchy, watery eyes ■ sneezing ■ itching of the nose or throat Do not use if you have ever had an allergic reaction to this product or any of its ingredients. Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose. When using this product do not take more than directed. Taking more than directed may cause drowsiness. Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away. If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away. Directions adults and children 6 years and over 1 tablet daily; not more than 1 tablet in 24 hours children under 6 years of age ask a doctor consumers with liver or kidney disease ask a doctor Other information ■ store between 20° to 25°C (68° to 77°F) Inactive ingredients

Unique Pharmaceutical Laboratories (A Div. of J. B. Chemicals & Manufactured by:

Manufactured for: Cabinet Health P.B.C. Brooklyn, NY 11222

#### For Repackaging Only

com starch, lactose monohydrate, magnesium stearate, povidone Questions or Comments? Call + 1-908-242-6108

Mfg. Lic. No. G/1430

100,000 Tablets 142292

#### LORATADINE

loratadine tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82725-0142	
Route of Administration	ORAL			

#### **Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN) **LORATADINE** 10 mg

Inactive Ingredients			
Ingredient Name	Strength		
STARCH, CORN (UNII: O8232NY3SJ)			
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
POVIDONE (UNII: FZ989GH94E)			

#### **Product Characteristics**

Color	white	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	P;10
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:82725- 0142-1	8000 in 1 BAG; Type 0: Not a Combination Product	10/01/2025		
2	NDC:82725- 0142-3	10000 in 1 BAG; Type 0: Not a Combination Product	10/01/2025		
3	NDC:82725- 0142-4	1 in 1 DRUM	10/01/2025		
3		96000 in 1 DRUM; Type 0: Not a Combination Product			
4	NDC:82725- 0142-2	1 in 1 DRUM	10/01/2025		
4		100000 in 1 BAG; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214684	10/01/2025	

# Labeler - Cabinet Health P.B.C. (117102391)

Revised: 10/2025 Cabinet Health P.B.C.